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EXAMPLE 32

N6-Benzoyl-5'-O-DMT-3'-O-(butylphthalimido)-adenosine

[0169] The title compound is prepared from 3'-O-(butyl-phthalimide)-N⁶-benzoyladenosine as per Example 22.

EXAMPLE 33

N6-Benzoyl-5'-O-DMT-3'-O-(butylphthalimido)-Adenosine-2'-O-(2-cyanoethyl-N,N-diisopropyl) phosphoramidite

[0170] The title compound is prepared from 3'-O-(butylphthalimide)-5'-O-DMT-N⁶-benzoyladenosine as per Example 24.

EXAMPLE 34

3'-O-(Pentylphthalimido)-adenosine

[0171] The title compound is prepared as per Example 21, using N-(5-bromopentyl)phthalimide. The crude material from the extraction is chromatographed on silica gel using CHCl₃/MeOH (95/5) to give a mixture of the 2' and 3' isomers. The 2' isomer is recrystallized from EtOH/MeOH 8/2. The mother liquor is rechromatographed on silica gel to afford the 3' isomer.

2'-O-(Pentylphthalimido)adenosine: M.P. 159-160 °C. Anal. Calcd. for C₂₃H₂₄N₆O₅: C, 57.26; H, 5.43; N, 17.42. Found: C, 57.03; H, 5.46; N, 17.33. 3'-O-(Pentylphthalimido)adenosine: ¹H NMR (DMSO-d₆) δ 5.87 (d, 1H, H-1').

EXAMPLE 35

N6-Benzoyl-3'-O-(pentylphthalimido)-adenosine

[0172] Benzoylation of 3'-O-(pentylphthalimido)adenosine is achieved as per the procedure of Example 22 to give the title compound.

EXAMPLE 36

N6-Benzoyl-5'-O-DMT-3'-O-(pentylphthalimido)-adenosine

[0173] The title compound is prepared from 3'-O-(pentyl-phthalimide)-N⁶-benzoyladenosine as per the procedure of Example 23. Chromatography on silica gel (ethylacetate, hexane, triethylamine), gives the title compound.

EXAMPLE 37

N6-Benzoyl-5'-O-DMT-3'-O-(pentylphthalimido)-adenosine-2'-O-(2-cyanoethyl-N,N-diisopropyl) phosphoramidite

[0174] The title compound is prepared from 3'-O-(pentyl-phthalimide)-5'-O-(DMT)-N⁶-benzovladenosine as per the procedure of Example 24 to give the title compound.

EXAMPLE 38

3'-O-(Propylphthalimido)uridine

[0175] A solution of uridine-tin complex (48.2 g, 115 mmol) in dry DMF (150 ml) and N-(3-

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bromopropyl)phthalimide (46 g, 172 mmol) was heated at 130 °C for 6 hrs. The crude product was chromatographed directly on silica gel CHCl₃/MeOH 95/5. The isomer ratio of the purified mixture was 2'/3' 81/19. The 2' isomer was recovered by crystallization from MeOH. The filtrate was rechromatographed on silica gel using CHCl₃CHCl₃/MeOH (95/5) gave the 3' isomer as a foam. 2'-*O*-(Propylphthalimide)uridine: Analytical sample recrystallized from MeOH, m.p. 165.5-166.5C, 'H NMR (200 MHZ, DMSO-d₆) δ 1.87 (m, 2H, CH₂), 3.49-3.65 (m, 4H, C₂H), 3.80-3.90 (m, 2H, C₃H₁C₄H), 4.09(m, 1H, C₂H), 5.07 (d, 1h, C₃OH), 5.16 (m, 1H, C₅OH), 5.64 (d, 1H, CH=), 7.84 (d, 1H, C₁·H), 7.92 (bs, 4H, Ar), 7.95 (d, 1H, CH=) and 11.33 (s, 1H, ArNH). Anal. C₂₀H₂₁N₃H₈, Calcd. C, 55.69; H, 4.91;, N, 9.74. Found, C, 55.75; H, 5.12; N, 10.01. 3'-*O*-(Propylphthalimide)uridine: 'H NMR (DMSO-d₆) δ 5.74 (d, 1H, H-1').

EXAMPLE 39

3'-O-(Aminopropyl)-uridine

[0176] The title compound is prepared as per the procedure of Example 25.

EXAMPLE 40

3'-O-[3-(N-trifluoroacetamido)propyl]-uridine

[0177] 3'-O-(Propylamino)uridine is treated as per the procedure of Example 26 to give the title compound.